High performance poly-Si TFTs fabricated by continuous-wave laser annealing of metal-induced lateral crystallised silicon films

C.-P. Chang and Y.S. Wu

In this process, amorphous silicon was first transformed to polycrystalline silicon (poly-Si) using a metal-induced lateral crystallisation (MILC) process, followed by annealing with a continuous-wave laser lateral ($\lambda \sim 532$ nm) crystallisation (CLC) with an output power of 3.8 W. MILC-CLC-TFT performed far superior to MILC-TFT. The mobility of the MILC-CLC-TFT was 293 cm²/Vs, which was much higher than that of MILC TFTs (54.8 cm²/Vs). In addition, MILC-CLC TFTs showed better device uniformity and reliability.

Introduction: Low-temperature polycrystalline silicon (LTPS) is very important for device applications such as solar cells and thin-film transistors (TFTs) [1]. Therefore, intensive studies have been carried out to reduce the crystallisation temperature of amorphous silicon (α -Si).

Among many methods, metal-induced lateral crystallisation (MILC) and excimer laser crystallisation (ELC) appear to be very promising methods [2–4]. MILC has the merits of low cost and uniform crystallisation over a large area. However, not all α -Si film was transformed to crystal Si [2, 3]. The ELC technique appears to be highly promising unfortunately their uniformity is inadequate and the surfaces of their poly-Si films are rough [4]. To improve the uniformity and performance of ELC-TFTs, many methods have been proposed [5–7]. The cost of the ELC system, however, is still high.

Recently, continuous-wave (CW) laser lateral crystallisation (CLC) of amorphous Si has been developed for LTPS TFT [8, 9]. Not only are the performances of CLC TFT better, but the manufacturing cost is lower than ELC TFTs. In this Letter, a new manufacturing method using post-annealing of MILC poly-Si TFTs with a CW laser (MILC-CLC) is proposed.

Experiment: The MILC process began with 4-inch quartz wafer substrates where wet oxide films of 500 nm were grown. A silane-based undoped amorphous silicon (α -Si) layer with a thickness of 100 nm was deposited using low-pressure chemical vapour deposition (LPCVD). The photoresist was patterned to form the desired Ni lines, and a 20 Å-thick Ni film was deposited on the α -Si. The samples were then dipped into acetone for 5 min to remove the photoresist. Samples were subsequently annealed at 540°C for 18 h to form the MILC poly-Si film. The unreacted Ni metal was removed by chemical etching. The MILC poly-Si films were then irradiated by a CW laser $(\lambda \sim 532 \text{ nm})$ with various output powers (2.5, 3.8, 5 W) in an air atmosphere to fabricate the MILC-CLC poly-Si. Reactive ion etching (RIE) was employed to form islands of poly-Si regions. Next, a 100 nm-thick oxide layer was deposited as the gate insulator by plasma-enhanced chemical vapour deposition (PECVD). A 200 nmthick poly-Si film was then deposited for gate electrodes by LPCVD. After defining the gate, self-aligned phosphorous ions were implanted to form the source/drain and gate. The dopant activation was performed at 600°C in N2 ambient for 24 h.

Results and discussions: Fig. 1a shows an SEM image of the Seccoetched MILC needle grains. Most of the grains were parallel to each other in the <111> direction [10]. The width of the needlelike grains was around 50 nm. Among these grains remained some uncrystallised α -Si regions, which had been etched away. To fabricate MILC-CLC poly-Si films, MILC films were irradiated using a CW laser with the scan direction parallel to the needlelike poly-Si grains. When the laser output power was 2.5 W, the sizes and shapes of the needle Si grains were similar to those of MILC poly-Si (Fig. 1a). This is because MILC-CLC films were in the amorphous-melting regime. Only the α -Si regions among Si grains were melted. When the output power reached 3.8 W, the width of the grains increased markedly from 50 nm to ${\sim}3~\mu\text{m},$ as shown in Fig. 1b. The large grains were only molten partially and served as the nuclei for growth. The width of these grains markedly increased to \sim 3 μ m owing to the geometrical coalescence of Si needle grains [11]. The grain boundary between grains disappears, resulting in the sudden development of a much larger grain. This coalescence is an important phenomenon for grains

having a strong preferred orientation (MILC needle grains had a strong preferred orientation <111>). In this study, the effect of the CW ($\lambda = 532$ nm) laser post-annealing was much better than that of the excimer laser. The width of the Si grains dramatically increased to ~3 µm (the grains ranged from 2.5 to 4 µm), while that of excimer laser post-annealing was only 600 nm [10]. When the laser power was 5 W, the width of the grains ranged from 3 to 12 µm. The uniformity of the grain size was poor. As a result, device performances were not uniform. To achieve high performance with good uniformity, the CW laser with an output power of 3.8 W was chosen to fabricate MILC-CLC TFTs.



Fig. 1 SEM images of MILC and MILC-CLC poly-Si grains a MILC b MILC-CLC



Fig. 2 Typical I_{DS} - V_{GS} transfer characteristics and field-effect motilities of MILC and MILC-CLC TFTs

Fig. 2 shows the transfer characteristics and field-effect mobility against the gate voltage of MILC-CLC and MILC TFTs. The measured and extracted key device parameters are summarised in Table 1. MILC-CLC TFTs exhibited field-effect mobility reaching 293 cm²/Vs, which was much higher than that of MILC TFTs. The subthreshold slope (SS) and V_{TH} of the MILC-CLC TFTs were 0.39 V/dec. and -4.54 V, which were superior to 1.42 V/dec. and 2.24 V of the MILC TFTs. The ON/OFF current ratios of the MILC-CLC and MILC poly-Si TFTs were 6.69×10^7 and 0.18×10^6 at $V_{DS} = 5$ V, respectively.

Table 1: Device characteristics of MILC and MILC-CLC TFTs

| Device parameters | MILC-CLC | MILC |
|---|----------------------|----------------------|
| Field-effect mobility (cm ² /Vs) | 293 | 54.8 |
| Subthreshold slope (V/dec.) | 0.39 | 1.42 |
| Threshold voltage V_{TH} (V) | -4.54 | 2.24 |
| ON/OFF current ratio | 6.69×10^{7} | 0.18×10^{6} |

As mentioned earlier, many intragrain defects and α -Si regions remained among MILC poly-Si grains. These defects trap charge carriers and degrade electric performance. MILC-CLC TFTs do not have these problems because, as presented in Fig. 1*b*, the width Si of MILC-CLC grains dramatically increased to 3 μ m. Most of these geometrical coalescence grains and their boundaries are parallel to the drain current (I_{ds}), reducing the impedance to carrier flow and thereby reducing the threshold voltage and greatly increasing the mobility and $I_{\rm on}/I_{\rm off}$ current ratio.

Twenty MILC-CLC TFTs were measured in μ_{FE} and V_{TH} to investigate device uniformity. The standard deviations of the μ_{FE} and V_{TH} are 7.46 and 0.165, respectively. As a result, small standard deviations of MILC-CLC TFTs indicate a fine uniformity owing to the CW laser annealing. The other important issue of MILC-CLC poly-Si TFTs is their reliability, which was examined under hot-carrier stress (HCS). Fig. 3 shows the field-effect mobility and threshold voltage variation against stress time, and the stress condition is $V_{D,stress} = 15 \text{ V}$, $V_{G,stress} = V_{GS} - V_{TH} = 15$ V for varied time duration. It is found that MILC-CLC TFTs also had a good reliability.



Fig. 3 Field-effect mobility and threshold voltage variation examined under hot-carrier stress

Conclusions: A high-performance LTPS TFT fabricated by MILC-CLC was investigated. In this process, amorphous silicon was first transformed to poly-Si using an MILC method, and then annealed using a continuous-wave laser. Laser-annealing with an output power of 3.8 W greatly increased the width of the needle grains from 50 nm to 3 µm by geometrical coalescence. MILC-CLC-TFT markedly outperformed that of the MILC-TFT because the MILC-CLC poly-Si film had much larger grains and fewer intragrain defects than the MILC poly-Si film. The MILC-CLC-TFT has a lower threshold voltage, smaller subthreshold slope and a higher on/off current ratio than the MILC-TFT. The mobility of the MILC-CLC-TFT was 293 cm²/Vs, which was much higher than that of MILC TFTs (54.8 cm²/Vs). Besides, MILC-CLC TFTs showed better device uniformity and reliability.

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